AMENDMENTS TO THE CLAIMS

- 1. (Currently Amended) A method of making an antibody molecule, the antibody containing an immunoglobulin heavy chain comprising a α 3 domain or a mu domain, the method comprising:
 - (a) Providing a nucleotide sequence encoding—the an immunoglobulin heavy chain molecule;
 - (b) Modifying the nucleotide sequence to form a modified nucleotide sequence, wherein the modifying is in the region of the nucleotide sequence encoding the C-terminus 18 amino acids of the immunoglobulin heavy chain molecule to remove, or reduce the effectiveness of, one or more vacuolar targeting signal of the encoded immunoglobulin heavy chain;
 - (c) Inserting the modified nucleotide sequence into a host cell; and
 - (d) Causing the host cell to express the modified nucleotide sequence to form a modified immunoglobulin heavy chain and secrete the modified immunoglobulin heavy chain from the host cell.

2-33. (Cancelled)

- 34. (**Previously Presented**) A method according to claim 1 wherein the immunoglobulin heavy chain molecule is IgA, IgM or an IgA/G hybrid.
- 35. (**Previously Presented**) A method according to claim 1 wherein the nucleotide sequence is modified by at least one of the modifications selected from the group consisting of
 - (i) one or more point mutations of the nucleotide sequence,
 - (ii) deleting one or more nucleotides,

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- (iii) adding one or more nucleotides and
- (iv) replacing one or more nucleotides with a synthetic nucleotide sequence.
- 36. (**Previously presented**) A method according to claim 35, wherein the synthetic nucleotide sequence encodes an amino acid sequence of general formula:

 $-(Xaa_1)_m C(Xaa_2)_n$

where: C

= a cysteine residue

Xaaı

independently any amino acid with the proviso that it is not from

I, L or forms a consecutive sequence X₁ X₂ X₃ V S X₄ (SEQ ID NO: 1)

where:

 $X_1 = N$, H or L

 $X_2 = V \text{ or } Y$

 $X_3 = S \text{ or } N$

 X_4 = aliphatic amino acid

Xaa₂ = independently any amino acid

m = at least 2

n = 0 to 5.

- 37. (Previously presented) A method according to claim 36, wherein Xaa_2 is Y and n = 1.
- 38. (Previously Presented) A method according claim 1, wherein nucleotides encoding the last 16 amino acids of the immunoglobulin heavy chain are deleted.
- 39. (Currently Amended) A method according to claim 1 wherein the resultant amino acid sequence at the C terminus of the immunoglobulin heavy chain has a formula selected from the group consisting of:
 - (a) SCMVGHEALPMNFTQKTIDRLSGKPACY (SEQ ID NO: 7),
 - (b) SCMVGHEALPMNFTQKTIDRLSGKPAAACY (SEQ ID NO: 8),
 - (c) SCMVGHEALPMNFTQKTIDRLSGKPHASTPEPDPVACY (SEQ ID NO: 9) and
 - (d) SCMVGHEALPMNFTQKTIDRLSGKPAAAAACY (SEQ ID NO: 69).
- 40. (**Previously Presented**) A method according to claim 1 wherein the nucleotide sequence of part (a) originally encoded the amino acid sequence:

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X₁ X₂ X₃ V S X₄ (SEQ ID NO: 1)

where: $X_1 = N$, H or L

 $X_2 = V \text{ or } Y$

 $X_3 = S \text{ or } N$

 X_4 = aliphatic amino acid.

- 41. (**Previously presented**) A method according to claim 40, wherein the amino acid sequence is: N V S V S V (SEQ ID NO: 2).
- 42. (**Previously Presented**) A method according to claim 1 wherein the nucleotide sequence of part (a) encoded L or I.
- 43. (**Currently Amended**) A method according to claim 42, wherein the modified nucleotide sequence encodes a modified amino acid selected from the group consisting of: (i) an isoleucine 3 amino acids from the C-terminus end of the immunoglobulin heavy chain,
- (ii) an isoleucine 10 amino acids from the C-terminus end of the immunoglobulin heavy chain and
- (iii) an isoleucine 3 amino acids from the C-terminus end of the immunoglobulin heavy chain and an isoleucine-to 2 amino acids from the C-terminus end of the immunoglobulin heavy chain.
- 44. (**Currently Amended**) A method according to claim 1, wherein the <u>method modified</u> nucleotide sequence is contained within a nucleotide sequence encoding the sequence:

PT X₁ X₂ X₃ V S X₄ X₅ X₆ X₇ X₈ X₉ X₁₀ X₁₁ X₁₂ C X₁₃ (SEQ ID NO: 5)

where: $X_1 = N$, H or L

 $X_2 = V \text{ or } Y$

 $X_3 = S \text{ or } N$

 X_4 = an aliphatic amino acid

 X_5 = an aliphatic amino acid

 $X_6 = M, V \text{ or } L$

 $X_7 = S \text{ or } A$

 $X_8 = E \text{ or } D$

 X_9 = any amino acid

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 $X_{10} = D, E, G \text{ or } A$

 $X_{11} = G \text{ or } S$

 $X_{12} = I, T, V, Z \text{ or } A$

 X_{13} = may or may not be present and, where present is A or $Y_{\underline{.}}$

- 45. **(Previously Presented)** A method of adding J-chain binding capability to the immunoglobulin heavy chain of an antibody comprising the steps of:
 - (a) providing a nucleotide encoding an immunoglobulin heavy chain;
 - (b) adding to that nucleotide a nucleotide sequence encoding a synthetic tail with the amino acid sequence:

$$-(Xaa_1)_m C(Xaa_2)_n$$

where: C = Cys

 Xaa_1 is independently any amino acid with the proviso that it is not I or L or forms a consecutive sequence $X_1 X_2 X_3 V S X_4$ (SEQ ID NO: 1) (where $X_1 = N$, H or L; $X_2 = N$)

V or Y;
$$X_3 = S$$
 or N; $X_4 =$ aliphatic amino acid)

 $Xaa_2 = independently any amino acid$

m = at least 2

n = 0 to 5; and

- (c) expressing the immunoglobulin nucleotide in a host cell to form an immunoglobulin heavy chain capable of binding J-chain.
- 46. **(Previously presented)** A method according to claim 1 wherein the host cell is a plant cell.
- 47. **(Previously presented)** A method according to claim 45 wherein the host cell is a plant cell.
- 48. (**Previously presented**) A method according to claim 46, wherein the plant cell is part of a transgenic plant.

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- 49. (**Previously presented**) A method according to claim 47, wherein the plant cell is part of a transgenic plant.
- 50. (**Previously presented**) A method according to claim 1 additionally comprising the step of isolating and purifying the antibody molecule.
- 51. (**Previously presented**) A method according to claim 45 additionally comprising the step of isolating and purifying the antibody molecule.
- 52. (**Previously Presented**) A method according to claim 50, wherein the antibody molecule is subjected to a protease digest to produce Fab or F(ab')₂ fragments.
- 53. (Previously presented) A method according to claim 51, wherein the antibody is subjected to a protease digest to for Fab or F(ab')₂ fragments.
- 54. (**Previously Presented**) An antibody containing an immunoglobulin heavy chain comprising an α 3 domain or a mu domain, the α 3 domain or mu domain lacking one or more targeting signals towards the C-terminal end.
- 55. (**Previously presented**) An antibody capable of binding J-chain comprising at its C-terminal end the sequence:

$$-(Xaa_1)_m C(Xaa_2)_n$$

where: $C = Cys$

 Xaa_1 is independently any amino acid with the proviso that it is not I or L or forms a consecutive sequence $X_1 X_2 X_3 V S X_4$ (SEQ ID NO: 1) (where $X_1 = N$, or

L;
$$X_2 = V$$
 or Y; $X_3 = S$ or N; $X_4 =$ aliphatic amino acid)

 $Xaa_2 = independently any amino acid$

$$m = at least 2$$

$$n = 0 \text{ to } 5$$

56. (Previously presented) An antibody according to claim 54 which does not contain the targeting signal: $X_1 X_2 X_3 V S X_4$ (SEQ ID NO: 1)

where:
$$X_1 = N$$
, H or L

$$X_2 = V \text{ or } Y$$

 $X_3 = S \text{ or } N$

 X_4 = aliphatic amino acid.

57. (Previously presented) An antibody according to claim 55 which does not contain the targeting signal: $X_1 X_2 X_3 V S X_4$ (SEQ ID NO: 1)

where: $X_1 = N$, H or L

 $X_2 = V \text{ or } Y$

 $X_3 = S \text{ or } N$

 X_4 = aliphatic amino acid.

- 58. (**Previously presented**) An antibody according to claim 56, wherein the targeting signal is N V S V S V (SEQ ID NO: 2).
- 59. (**Previously presented**) An antibody according to claim 57, wherein the targeting signal is N V S V S V (SEQ ID NO: 2).
- 60. (**Previously presented**) An antibody according to claim 54 which contains one or no isoleucine or leucine amino acids within the last 18 amino acids at the C-terminus of the heavy chain of the antibody.
- 61. (Previously presented) An antibody according to claim 55 which contains one or no isoleucine or leucine amino acids within the last 18 amino acids at the C-terminus of the heavy chain of the antibody.
- 62. (**Previously presented**) An antibody according to claim 54 comprising at the C-terminus end of the heavy chain of antibody, the sequence:

$$-(Xaa_1)_m C(Xaa_2)_n$$

where: C = cysteine residue

 Xaa_1 = independently any amino acid with the proviso that it is not I or L or forms a consecutive sequence $X_1 X_2 X_3 V S X_4$ (SEQ ID NO: 2)

where:
$$X_1 = N$$
, H or L
 $X_2 = V$ or Y

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$$X_3 = S \text{ or } N$$

 X_4 = aliphatic amino acid

Xaa₂ = independently any amino acid

m = at least 2

n = 0 to 5.

63. **(Previously presented)** An antibody according to claim 55 comprising at the C-terminus end of the heavy chain of antibody, the sequence:

 $-(Xaa_1)_m C(Xaa_2)_n$

where: C = cysteine residue

 Xaa_1 = independently any amino acid with the proviso that it is not I or L or forms a consecutive sequence $X_1 X_2 X_3 V S X_4$ (SEQ ID NO: 2)

where: $X_1 = N$, H or L

 $X_2 = V \text{ or } Y$

 $X_3 = S \text{ or } N$

 X_4 = aliphatic amino acid

Xaa₂ = independently any amino acid

m = at least 2

n = 0 to 5.

- 64. (Previously presented) An antibody according to claim 54 in which at least two, preferably two to four, glycine or alanine residues are present downstream of a C-terminal targeting sequence
- 65. (Previously presented) An antibody according to claim 55 in which at least two, preferably two to four, glycine or alanine residues are present downstream of a C-terminal targeting sequence
- 66. (**Previously presented**) An antibody according to claim 54 in which at least the terminal amino acid residue of a C-terminal targeting sequence is replaced by at least two, preferably two to four, glycine or alanine residues.

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67. (Previously presented) An antibody according to claim 55 in which at least the terminal

amino acid residue of a C-terminal targeting sequence is replaced by at least two, preferably two

to four, glycine or alanine residues.

68. (Previously presented) A method of treating a disease by administering an antibody

according to claim 54 to a patient.

69. (**Previously presented**) A method of treating a disease by administering an antibody

according to claim 55 to a patient.

70. (**Previously presented**) A method of prophylaxis, comprising administering an antibody

according to claim 54 to a person or animal.

71. (**Previously presented**) A method of prophylaxis, comprising administering an antibody

according to claim 55 to a person or animal.

72. (Previously presented) A vector comprising a nucleotide sequence encoding an antibody

according to claim 54.

73. (Previously presented) A vector comprising a nucleotide sequence encoding an antibody

according to claim 55.

74. (**Previously presented**) A host cell comprising a nucleotide sequence encoding antibody

according to claim 54.

75. (Previously presented) A host cell comprising a nucleotide sequence encoding antibody

according to claim 55.

76. (Previously presented) A host cell comprising a vector according to claim 72.

77. (**Previously presented**) A host cell comprising a vector according to claim 73.

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- 78. (**Previously presented**) A transgenic plant comprising a nucleotide encoding an antibody according to claim 54.
- 79. (**Previously presented**) A transgenic plant comprising a nucleotide encoding an antibody according claim 55.
- 80. (Previously presented) An immunoassay comprising an antibody as defined in claim 54.
- 81. (Previously presented) An immunoassay comprising an antibody as defined in claim 55.
- 82. (**Previously Presented**) The method of claim 1, further comprising adding to the nucleotide sequence encoding the immunoglobulin heavy chain a nucleotide sequence encoding a synthetic tail with the amino acid sequence $-(Xaa_1)_m C(Xaa_2)_n$, wherein:
 - C = Cys
 - Xaa₁ is independently any amino acid with the proviso that it is not I or L or forms a consecutive sequence $X_1 X_2 X_3 V S X_4$ (where $X_1 = N$, H or L; $X_2 = V$ or Y; $X_3 = S$ or N; $X_4 =$ aliphatic amino acid)
 - Xaa₂ = independently any amino acid
 - -m = at least 2
 - n = 0 to 5; and

wherein said synthetic tail adds J-chain binding capability to the heavy chain of the immunoglobulin.

- 83. (**Previously presented**) A method according to claim 82 wherein the host cell is a plant cell.
- 84. (**Previously presented**) A method according to claim 83, wherein the plant cell is part of a transgenic plant.
- 85. (**Previously presented**) A method according to claim 82 additionally comprising the step of isolating and purifying the antibody molecule.

- 86. (**Previously Presented**) A method according to claim 85, wherein the antibody molecule is subjected to a protease digest to produce Fab or F(ab')₂ fragments.
- 87. (Currently Amended) The method according to claim 44, wherein at least one of X_1 - X_{13} is a member selected from the group consisting of:

$$X_1 = N$$

$$X_2 = V$$

$$X_4 = V \text{ or } L$$

$$X_5 = I, V \text{ or } L$$

$$X_6 = M$$

$$X_9 = G$$
, V, A or T

$$X_{10} = D$$

$$X_{11} = G$$

$$X_{12} = I$$
 or T .